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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO. CONFIRMATION NO.		
10/564,356	01/12/2006	Shin Yazawa	283520US0PCT	7829	
22850 7590 05/10/2007 OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT, P.C. 1940 DUKE STREET ALEXANDRIA, VA 22314			EXAMINER		
			PESELEV, ELLI		
ALEXANDRIA	A, VA 22314		ART UNIT PAPER NUMBER		
			1623		
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			NOTIFICATION DATE	DELIVERY MODE	
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

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	App	lication No.	Applicant(s)			
Office Action Summary		564,356	YAZAWA ET AL.			
		miner	Art Unit			
	Elli f	Peselev	1623			
The MAILING DATE of this com Period for Reply	The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply					
A SHORTENED STATUTORY PERIOD WHICHEVER IS LONGER, FROM THE Extensions of time may be available under the provafter SIX (6) MONTHS from the mailing date of this If NO period for reply is specified above, the maxim Failure to reply within the set or extended period for Any reply received by the Office later than three meanned patent term adjustment. See 37 CFR 1.70	HE MAILING DATE (risions of 37 CFR 1.136(a). In a communication. num statutory period will apply r reply will, by statute, cause onths after the mailing date of	OF THIS COMMUNICATION In no event, however, may a reply be tire of and will expire SIX (6) MONTHS from the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).			
Status						
 Responsive to communication(s This action is FINAL. Since this application is in cond closed in accordance with the p 	2b) ☐ This actio	n is non-final. cept for formal matters, pro				
Disposition of Claims						
4) ☑ Claim(s) <u>5-53</u> is/are pending in 4a) Of the above claim(s) 5) ☐ Claim(s) is/are allowed. 6) ☑ Claim(s) <u>5-53</u> is/are rejected. 7) ☐ Claim(s) is/are objected is are subject to re	is/are withdrawn fro					
Application Papers						
9) The specification is objected to to 10) The drawing(s) filed on is Applicant may not request that any Replacement drawing sheet(s) incli 11) The oath or declaration is object	/are: a) ☐ accepted objection to the drawinuding the correction is a	g(s) be held in abeyance. Sec required if the drawing(s) is ob	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).			
Priority under 35 U.S.C. § 119						
	of: ority documents have ority documents have bies of the priority do national Bureau (PC)	e been received. e been received in Application cuments have been receive	on No ed in this National Stage			
Attachment(s)						
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Reviolation Disclosure Statement(s) (PTO/SB Paper No(s)/Mail Date		4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal Pa	ite			

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The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 5-31 are rejected under 35 U.S.C. 103(a) as being unpatentable over the Japanese Patent (11-60592) or the Japanese Patent (200-191685) in view of Endo et al (U.S. Patent No. 6,569,464).

Each of the Japanese patents discloses cholestanol glycosides useful as anticancer agents but does not disclose said agents in liposome formulation. However, Endo et al disclose that "liposomes encapsulating drugs have been actively developed for the purposes of stabilization of unstable drugs, slow release of drugs in living bodies and targeting of drugs to lesion sites" (column 1, lines 15-25). Endo et al further disclose liposomal compositions comprising a wide variety of drugs, phospholipids and a positive-charge-providing substance such as cholesterol, aliphatic amines and fatty

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acids such as stearylamine and oleic acid (column 2, lines 30-56, column 4 and column 5, lines 1-15). Endo et al also teach that the drugs contained in the liposomes disclosed are not limited and may either hydrophilic or lipophilic (column 3, lines 56-59). Therefore, a person having ordinary skill in the art at the time the claimed invention was to made to prepare formulations of drugs disclosed by the Japanese Patents in liposomal formulations as disclosed by Endo et al in order to improve stabilization of said drugs and to slow release said drugs.

Claims 32-42 are rejected under 35 U.S.C. 103(a) as being unpatentable over Sarkar et al (The Journal of Biological Chemistry, vol. 272, vol. 41, pages 25608-25616, 1997) in view of Endo et al (U.S. Patent No. 5,569,464).

Sarkar et al disclose naphthalene glycosides drugs but do not disclose said compounds in liposome form. However, since Endo et al disclose the advantage of preparing a wide variety of drugs in liposomal form as discussed above, a person having ordinary skill in the art at the time the claimed invention was made would have been motivated to prepare drugs disclosed by Sakar et al in liposomal formulations.

Claims 43-53 are rejected under 35 U.S.C. 103(a) as being unpatentable over Akimoto et al (U.S. Patent No. 5,849,716) in view of Endo et al (U.S. Patent No. 5,569,464).

Akimoto et al disclose ceramide glycosides drugs having anti-tumor activity (column 1, lines 45-67 and column 2, lines 1-40) but do not disclose said drugs in liposome formulation. However, since Endo et al disclose the advantage of preparing a wide variety of compounds in liposomal formulation as discussed above, a person

having ordinary skill in the art at the time of the claimed invention would have been motivated to prepare said drugs in liposomal form.

Applicant's arguments filed March 5, 2007 have been fully considered but they are not persuasive.

Applicants contend that the Japanese Patents and Sarkar et al do not disclose incorporating said compounds into a liposomal composition. This argument has not been found persuasive because the advantage of incorporating drugs into liposomal compositions was well known in the art at the time the claimed invention was made as disclosed by Endo et al. Applicants further contend that Sarkar et al disclose utilizing a naphthalenyl glycoside as an anti-inflammatory agent and not an anti-tumor agent. This argument has not feen found persuasive since the present claims are directed to compositions and not to methods of use. In response to applicant's argument that Sarkar et al do not disclose the use of naphthalenyll glycosides as anti-tumor agents, a recitation of the intended use of the claimed invention must result in a structural difference between the claimed invention and the prior art in order to patentably distinguish the claimed invention from the prior art. If the prior art structure is capable of performing the intended use, then it meets the claim.

Applicants also contend that a skilled artisan would reasonably expect that the cholestanyl glycosides of the present invention would intrinsically possess sufficient affinity to cellular membranes without the aid of a liposomal carrier. This argument has not been found persuasive because Endo et al disclose the advantage of liposome

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formulations other than he affinity to cellular membranes such as slow release of drugs and targeting of drugs (column 1, lines 15-25).

Applicants also contend that inventor have unexpectedly discovered that by incorporating the cholestanyl glycoside anticancer agents into the liposomal carrier, the liposomal composition of the present invention exhibited enhanced antitumor efficacy as shown in Figures 1 and 2. This argument has not been found persuasive since the claims have not been limited to the compounds shown in Figures 1 and 2 i.e. the data shown in Figures 1 and 2 is not commensurate in scope with the claimed invention.

Applicants also contend that Endo et al disclose incorporating a carcinostatic active agent into a liposomal composition, said carcinostatic agent is present in a laundry list of more than twenty widely diversified types of active gents and that there is a lack of motivation to pick and chose agents therefrom. Applicants also contend that the anti-cancer agents disclosed by Endo et al are vastly different in chemical structure in comparison with the claimed compounds. This argument has not been found persuasive since Endo et al disclose incorporation into liposomes not only of anti-cancer drugs but a wide variety of other drugs and teach that the "drugs to be contained in the liposomes of the present invention are not particularly limited and may be either hydrophilic or lipophilic" (column 3, lines 56-59). Endo et al also disclose incorporating into liposomes hormonal preparations (column 4, lines 35-40) which have structurally similar to the cholestanol.

Therefore, the claimed compositions are still deemed prima facie obvious over the cited prior art.

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Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Elli Peselev whose telephone number is (571) 272-0659. The examiner can normally be reached on 8.00-4.30.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Shaojia Jiang can be reached on (571) 272-0627. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Elli Peselev

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